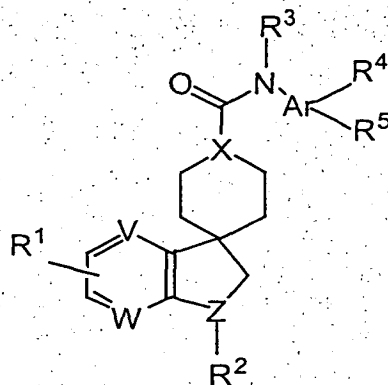


WHAT IS CLAIMED IS:

1. A compound of structural formula I:



I

or a pharmaceutically acceptable salt thereof, wherein;

V, W, X and Z are independently selected from CH and N;

R¹ is H, C₁₋₃ alkyl, C₁₋₃ alkoxy, F, or Cl;

R² is S(O)_n R⁶, COR⁶ or CHO, wherein

n is 0, 1 or 2; and

R⁶ is N(R³)₂ or C₁₋₃ alkyl;

R³ is independently H or C₁₋₃ alkyl;

Ar is aryl or heteroaryl;

R⁴ and R⁵ are independently selected from:

(1) hydrogen,

(2) aryl, either unsubstituted or substituted with

(a) halo

(b) C₁₋₃ alkoxy,

(c)-N(C₁₋₃ alkyl)₂,

(d) C₂₋₄ alkanoyl, or

(e) aryl;

(3) nitro,

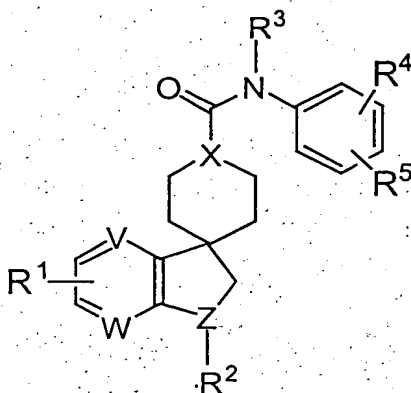
(4) C₁₋₅ alkyl,

(5) C₁₋₅ alkoxy,

(6) hydroxy-C₁₋₃ alkyl,

- (7) carboxy,
 (8) halo,
 (9) C₁₋₅ alkylthio,
 (10) C₁₋₅ alkoxy, carbonyl,
 (11) pyridyl, carbonyl,
 (12) benzoyl,
 (13) phenyl-C₁₋₃ alkoxy,
 (14) pyridyl, either unsubstituted or substituted with
 C₁₋₃ alkyl or C₁₋₃ alkoxy,
 (15) C₃₋₆ cycloalkyl,
 (16) oxazolyl,
 (17) thiazolyl,
 (18) triazolyl,
 (19) phenoxy or
 (20) C₂₋₆ alkanoyl.

2. The compound of Claim 1 wherein Ar is phenyl, of structural formula I(a)

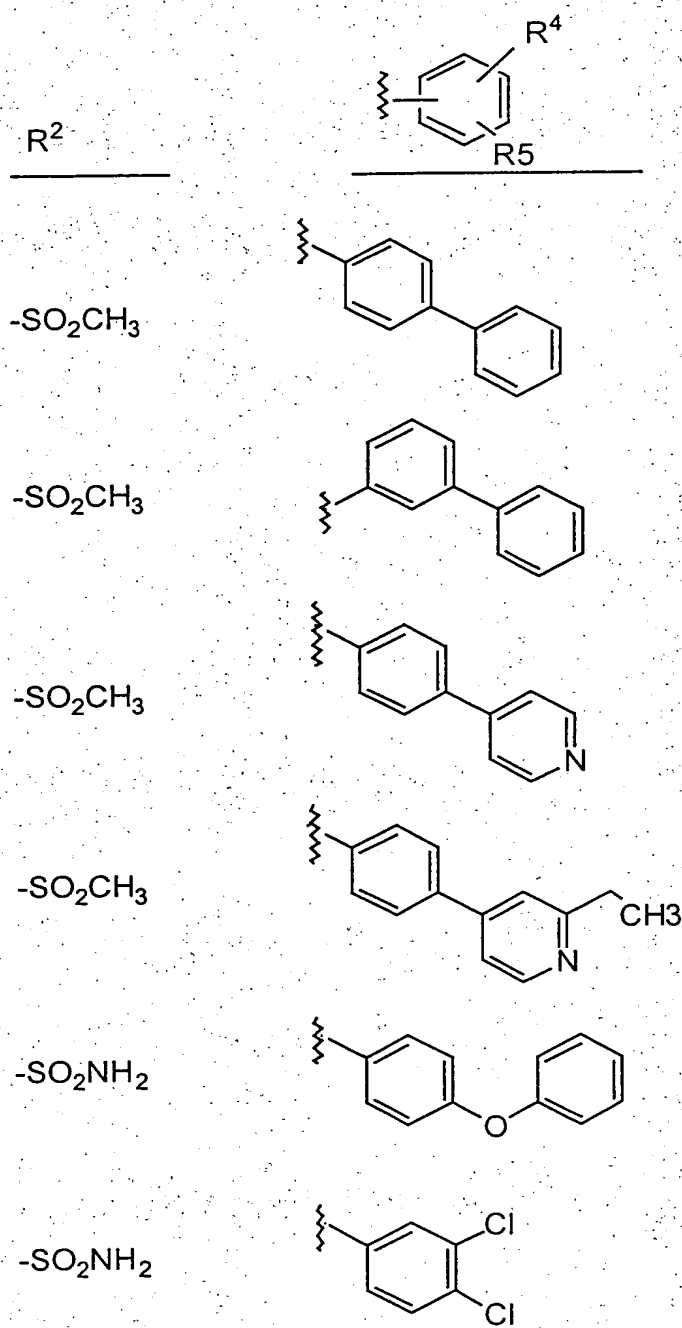


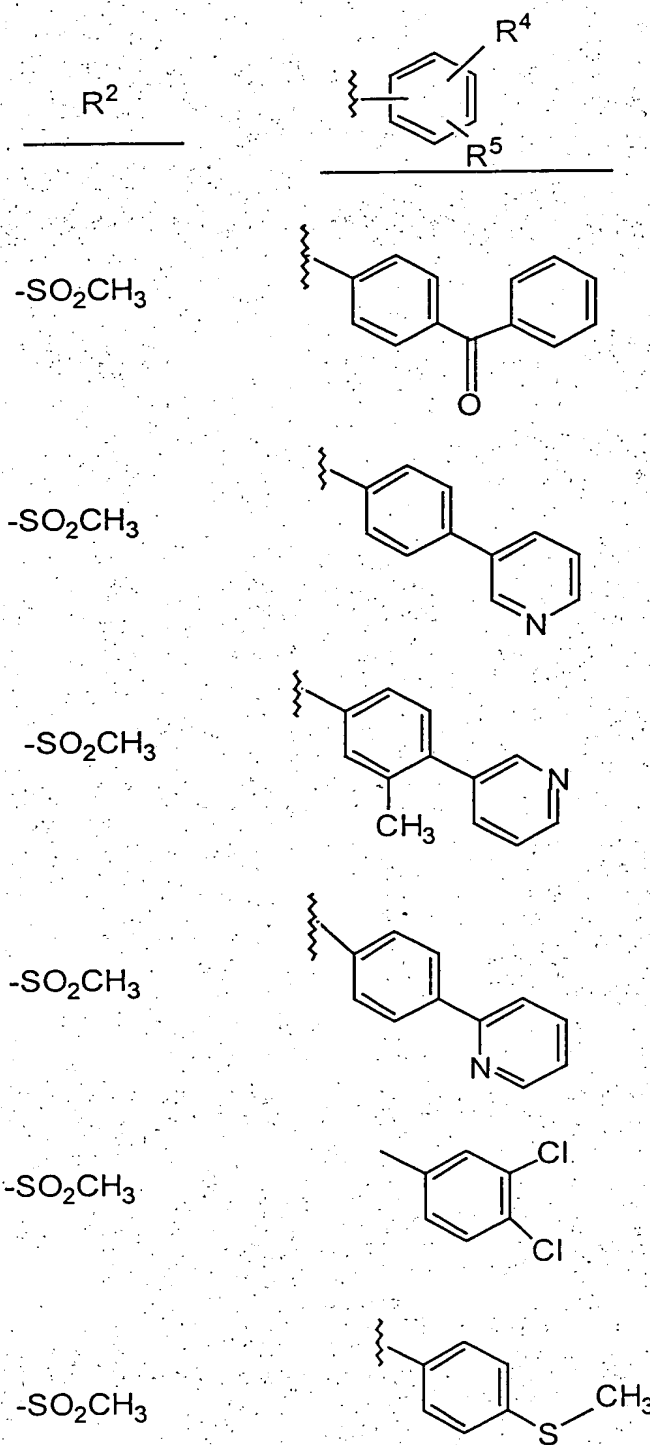
I(a)

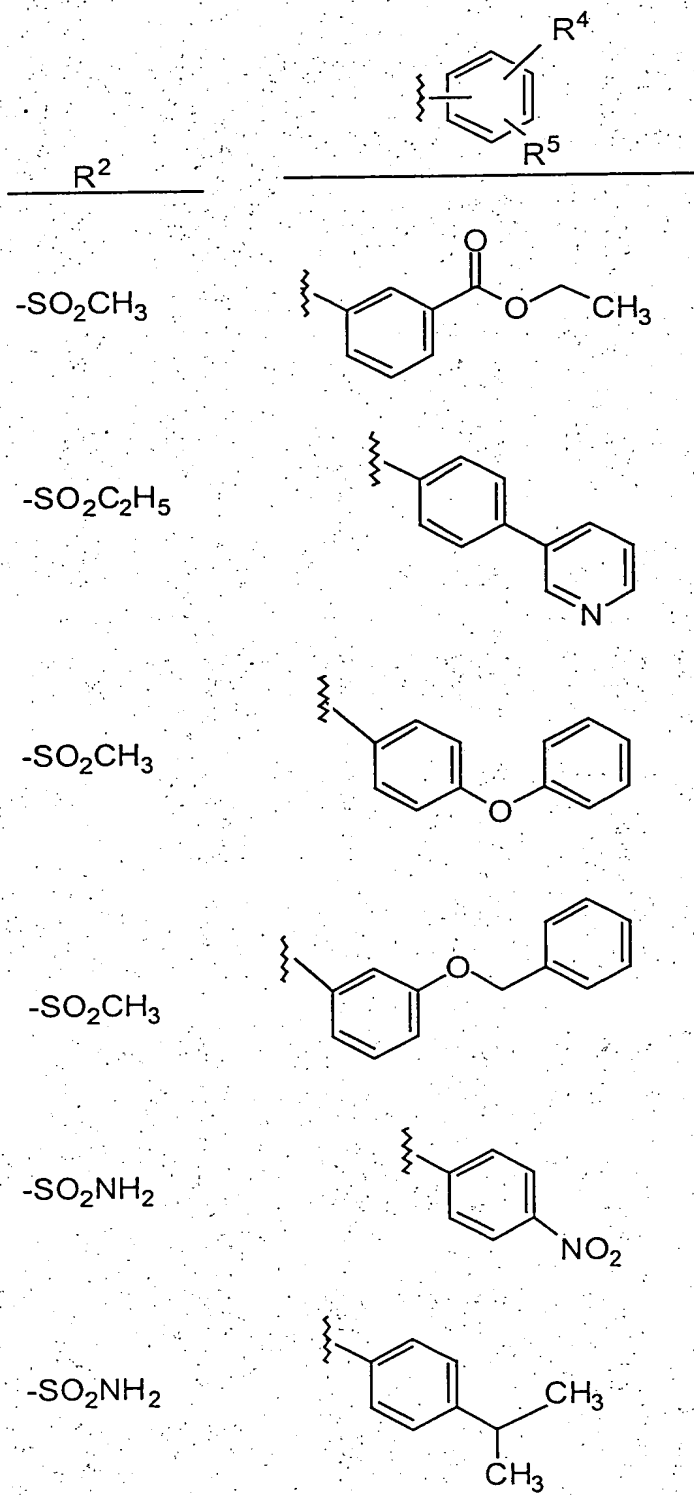
or a pharmaceutically acceptable salt thereof.

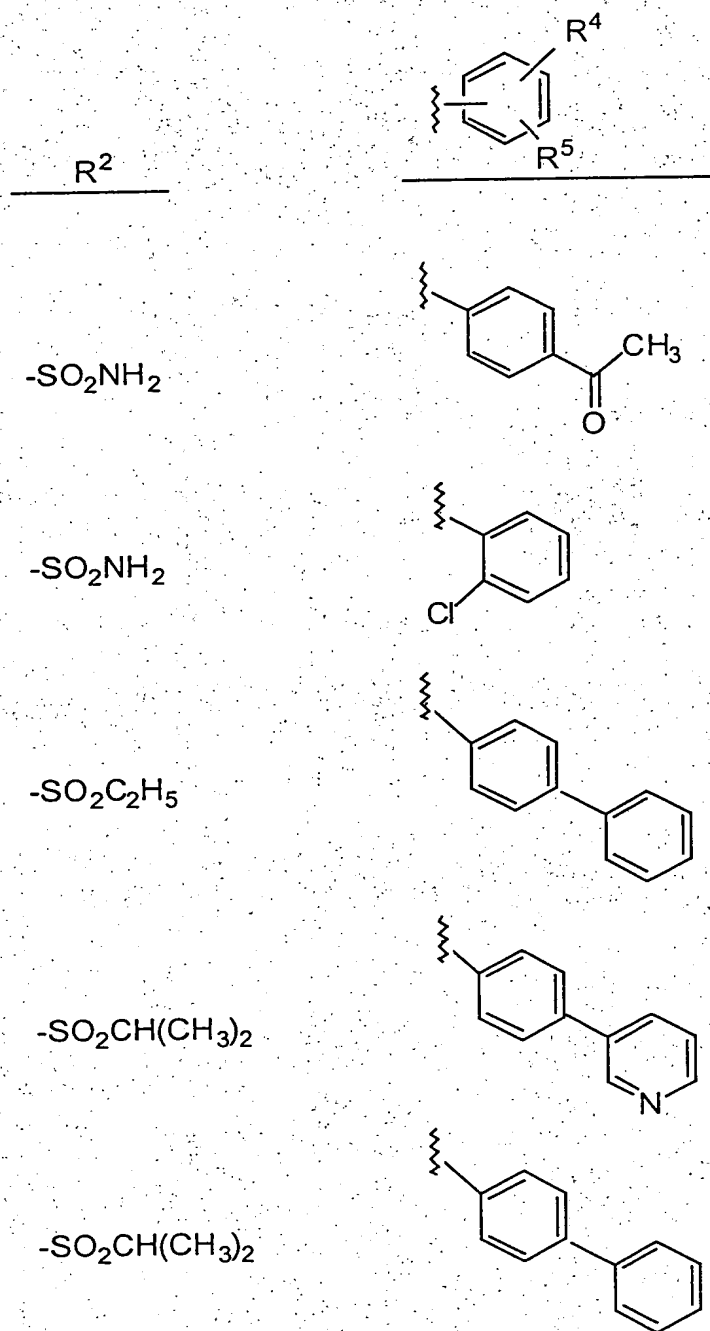
3. The compound of Claim 2 wherein X and Z are both nitrogen and V and W are both -CH=.

4. The compound of Claim 3 wherein R² is -SO₂(C₁₋₃ alkyl) or SO₂NH₂.
5. The compound of Claim 4 wherein R⁴ and R⁵ are independently selected from: phenyl, pyridyl, benzoyl, halophenyl, phenoxy, C₁₋₅ alkylpyridyl, benzhydryl, phenyl-C₁₋₃ alkoxy, NO₂, C₂₋₄ alkanoyl, halo, C₁₋₅ alkoxy, C₁₋₃ alkoxycarbonyl, C₁₋₅ alkylthio, triazolyl, carboxy, hydrogen, C₁₋₅ alkyl, pyridylcarbonyl, and C₁₋₃ alkoxyphenyl.
6. The compound of Claim 5 or a pharmaceutically acceptable salt thereof selected from those depicted in the following Table:

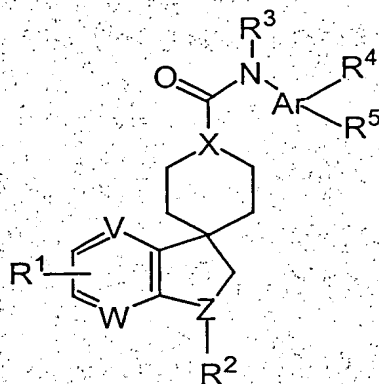








7. The compound of Claim 1 wherein Ar is a 5- or 6-membered heteroaryl having, besides carbon atoms, 1 to 3 hetero atoms selected from N, O or S as atoms constituting the ring, or benzo- or pyrido- fused versions thereof of structural formula I(b);



I(b)

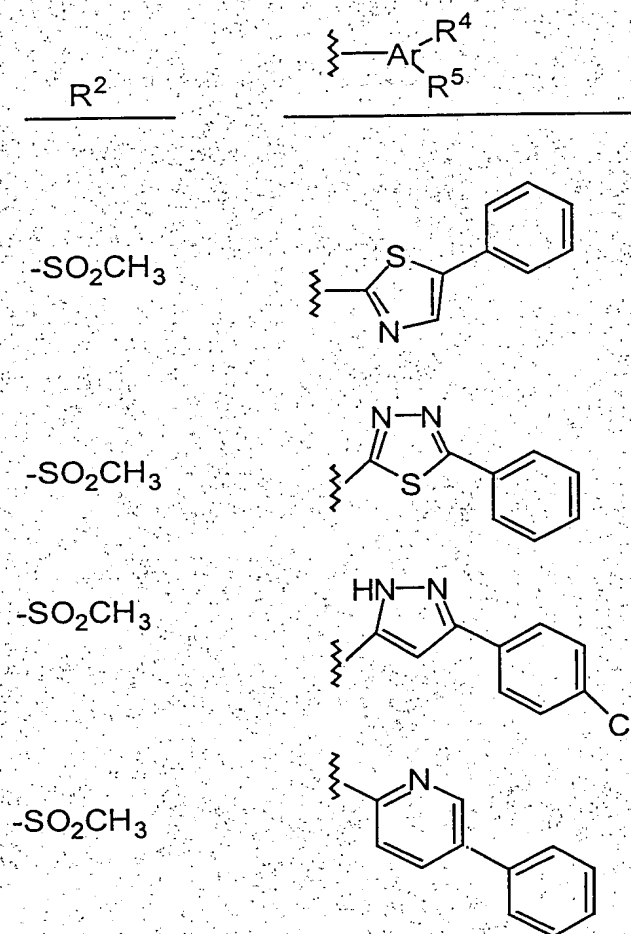
or a pharmaceutically acceptable salt thereof.

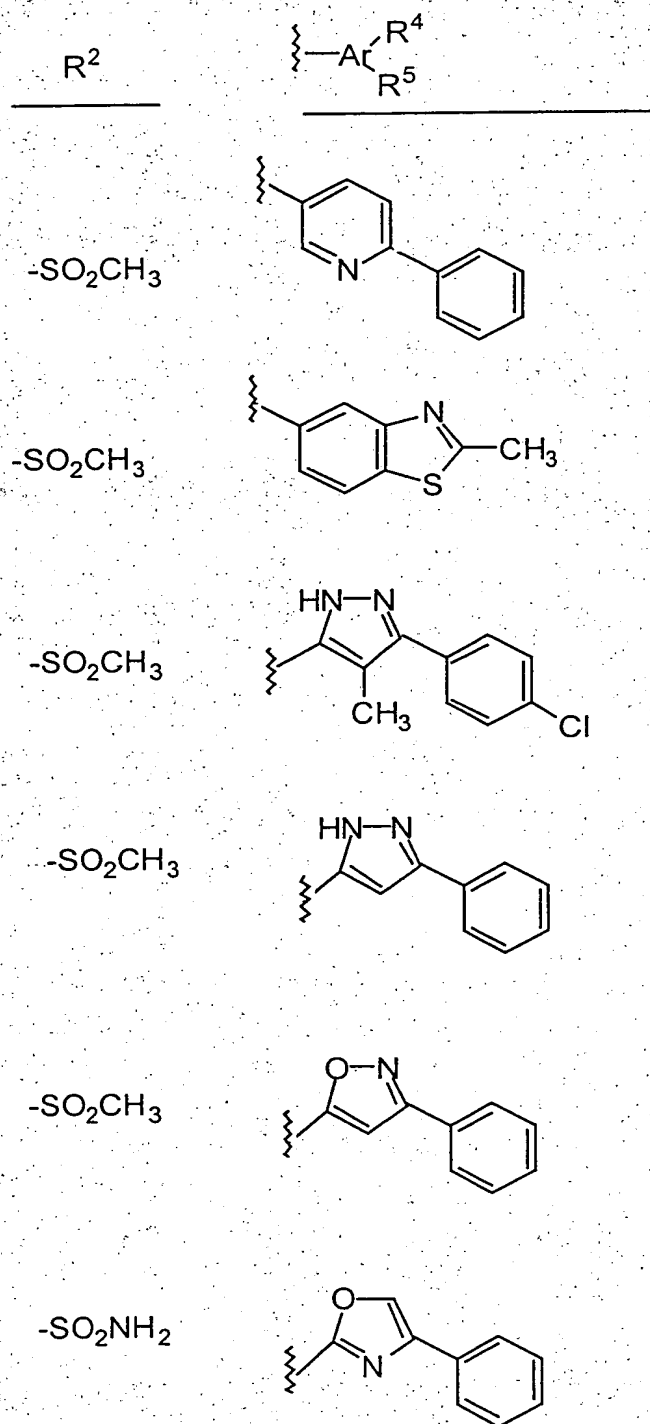
8. The compound of Claim 7 wherein X and Z are both nitrogen and V and W are both -CH=.

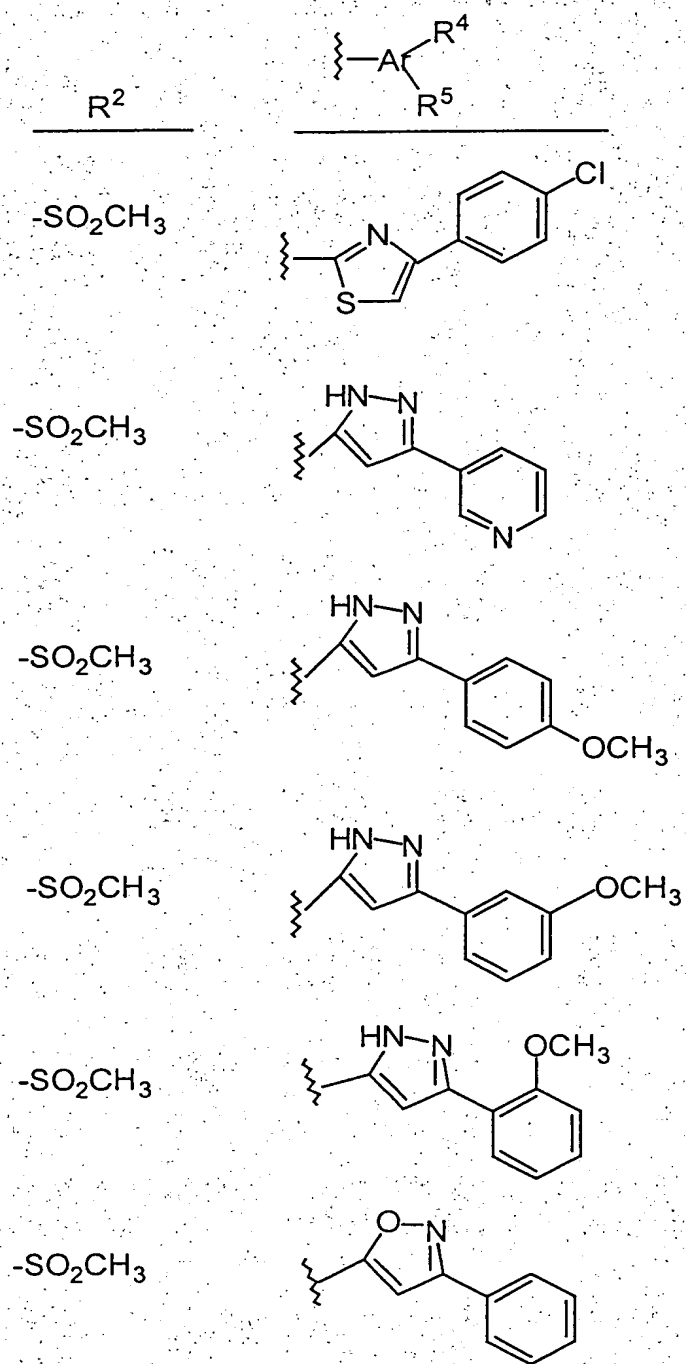
9. The compound of Claim 8 wherein R² is -SO₂(C₁₋₃ alkyl) or -SO₂N(C₁₋₃ alkyl)₂.

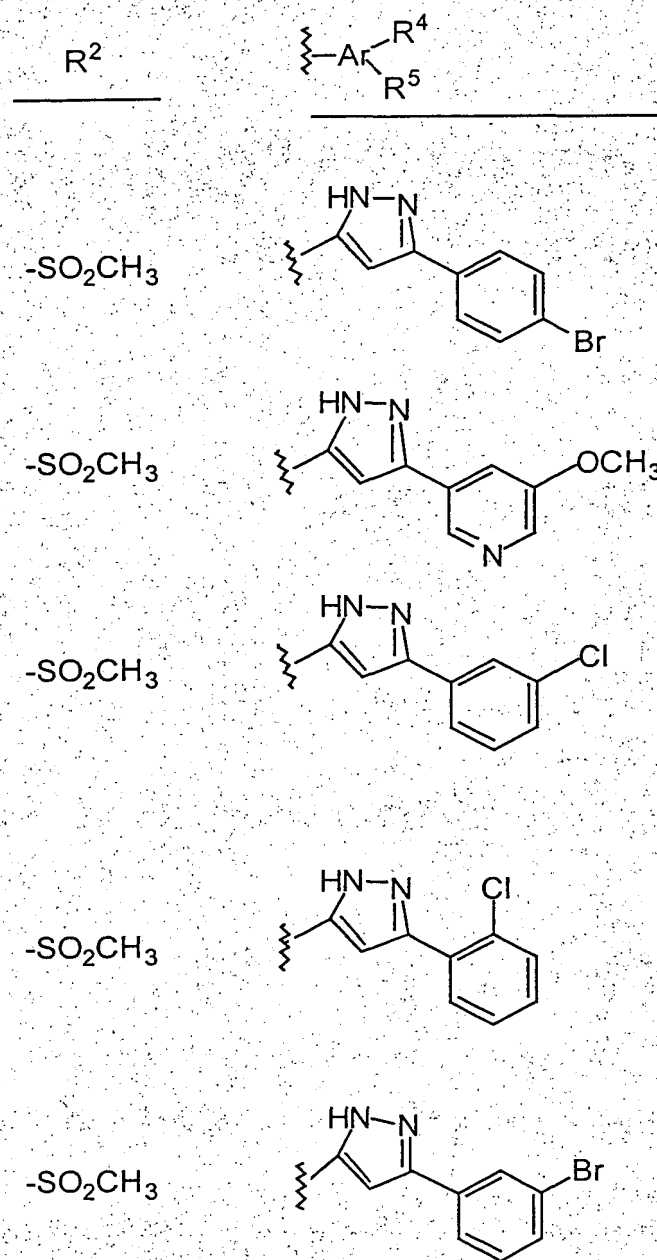
10. The compound of Claim 9 wherein the heteroaryl group, Ar, is selected from: thiazolyl, thiadiazolyl, pyrazolyl, pyridyl, benzothiazolyl, oxazolyl, pyridothiazolyl, benzoxazolyl, quinolyl, pyrazinyl, thienyl, isoxazolyl, pyrimidinyl, benzimidazolyl, oxadiazolyl and imidazolyl.

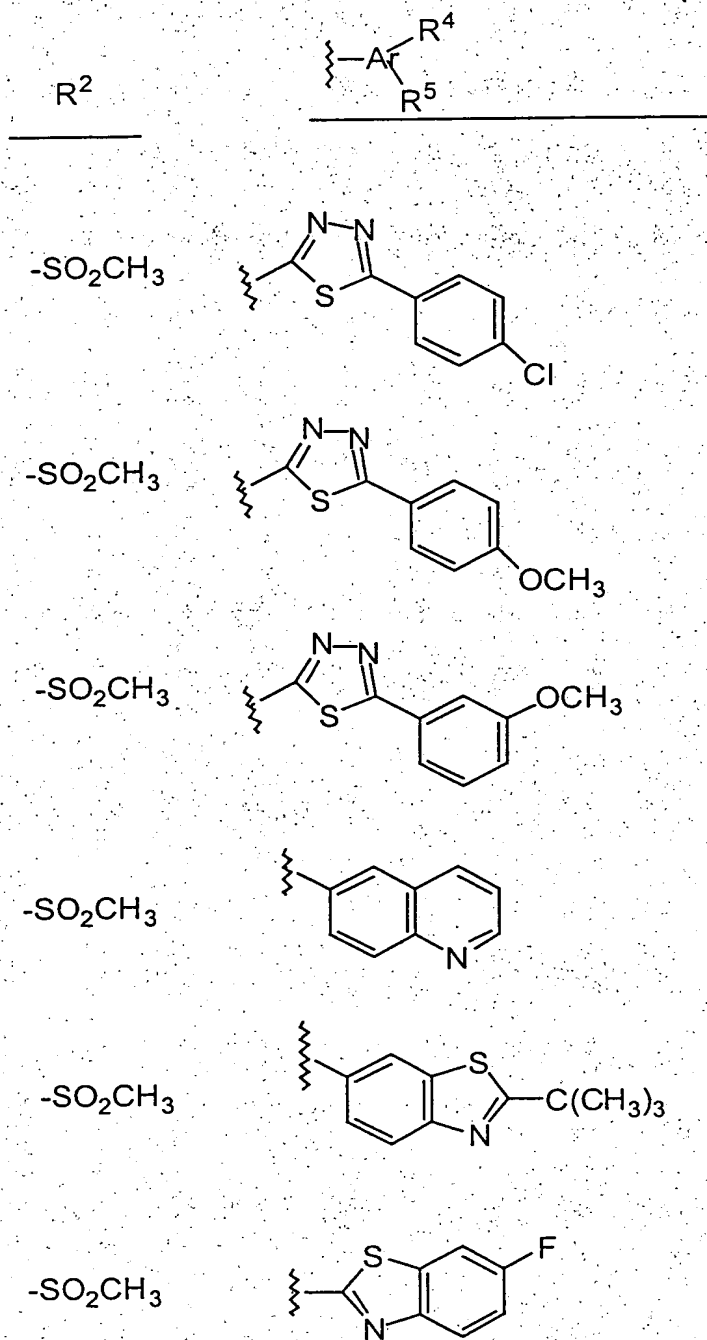
11. The compound of Claim 10, or a pharmaceutically acceptable salt thereof, selected from those depicted in the following Table:

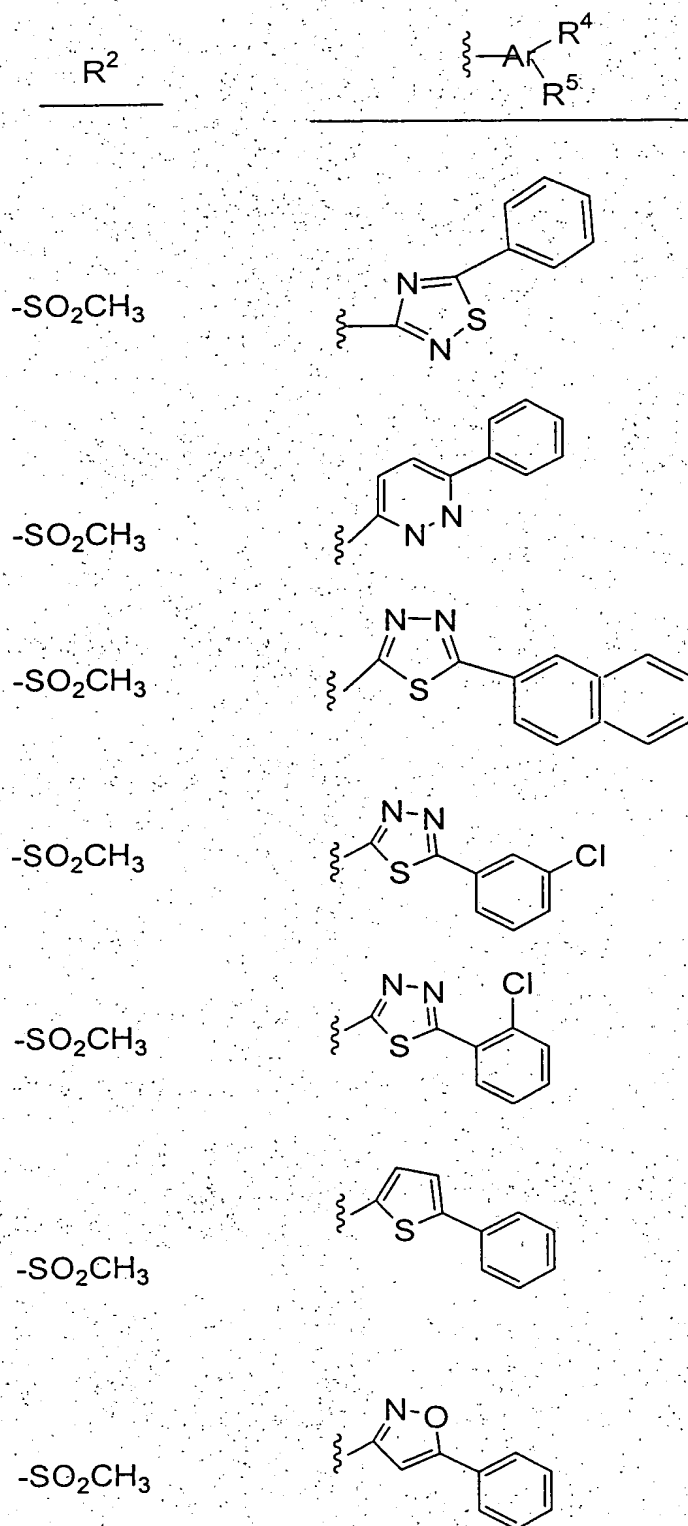


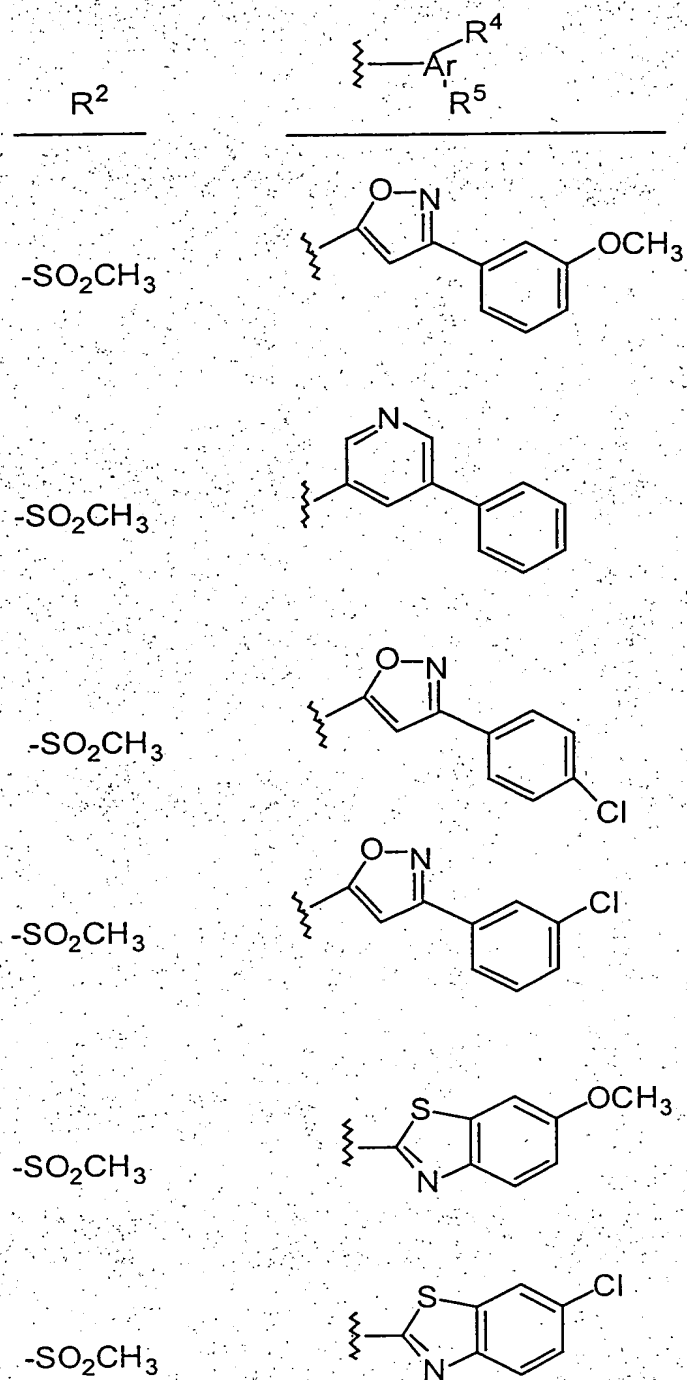


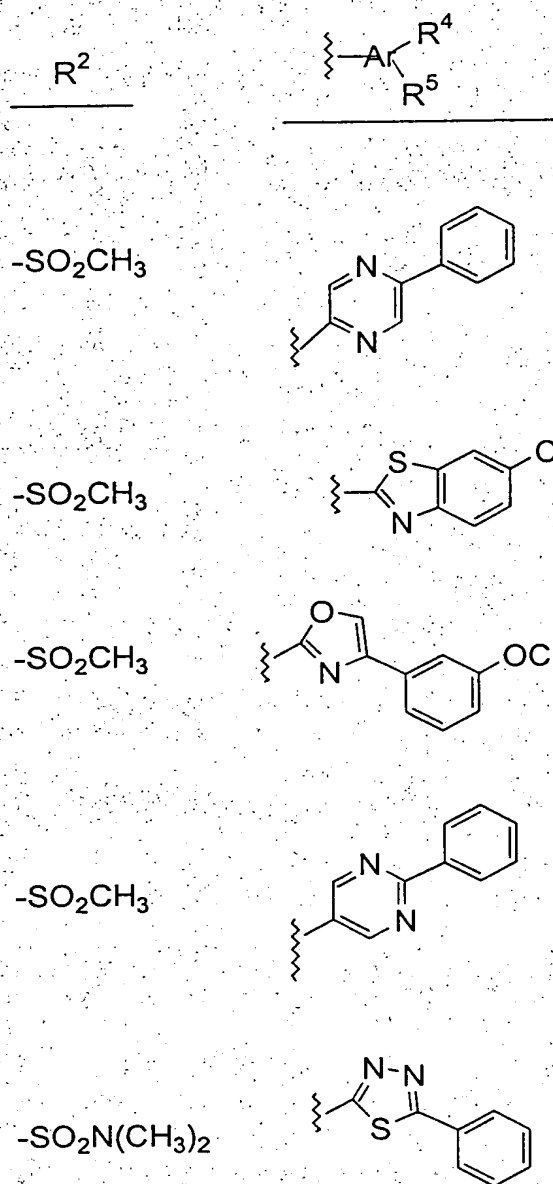


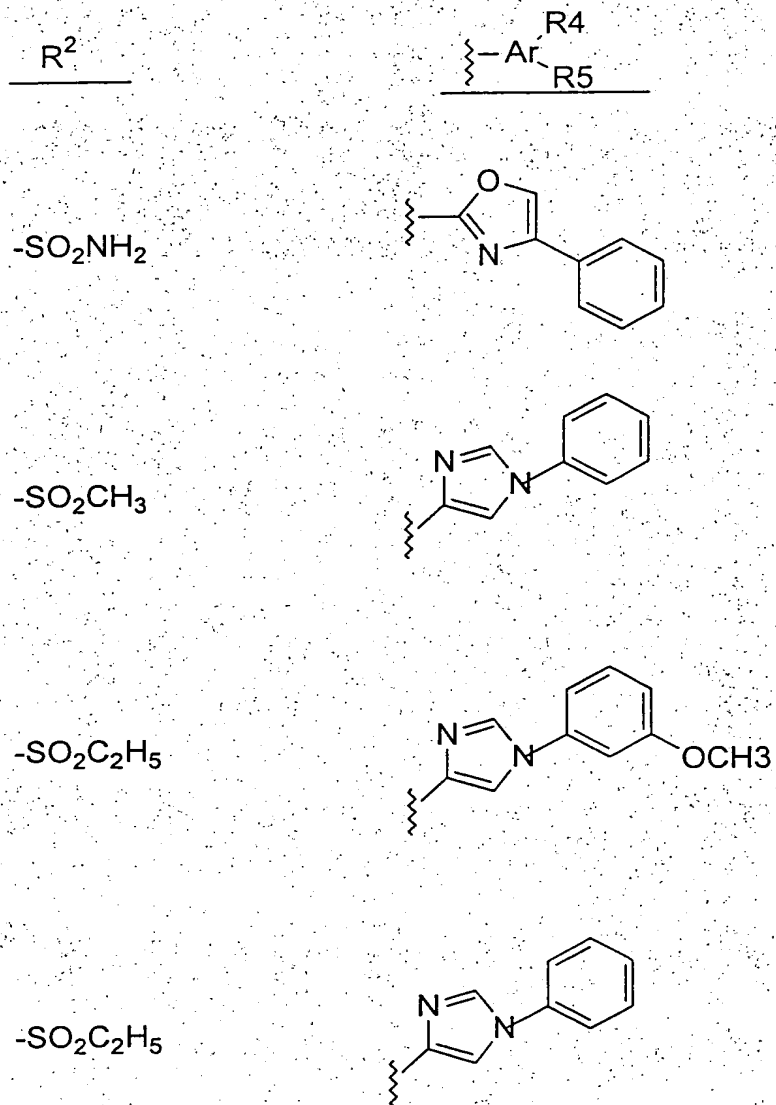


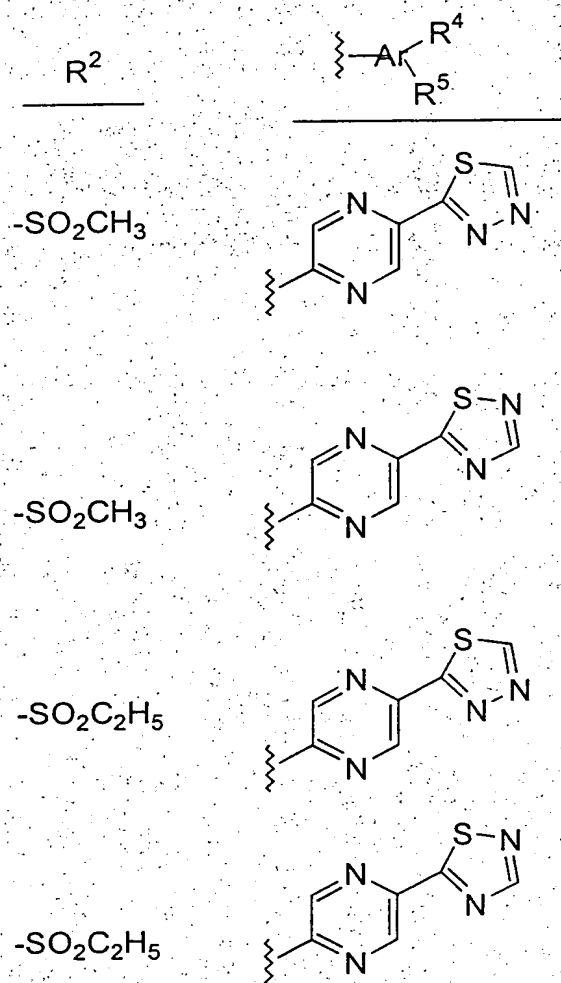




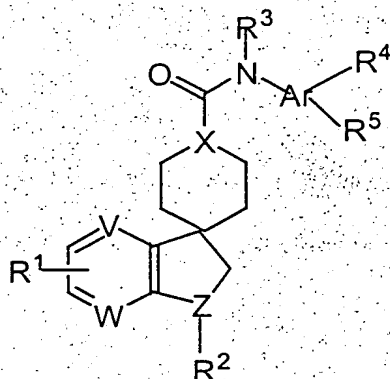








12. The compound of Claim 1 wherein one of X and Z is N and the other is -CH= of structural formula I(c):



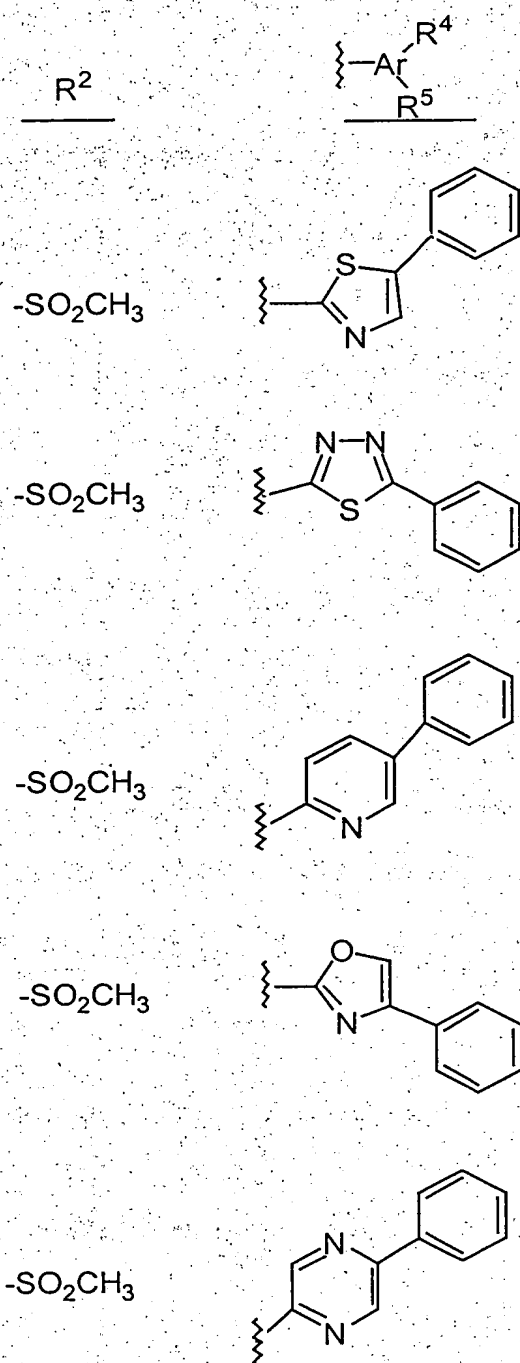
I(c)

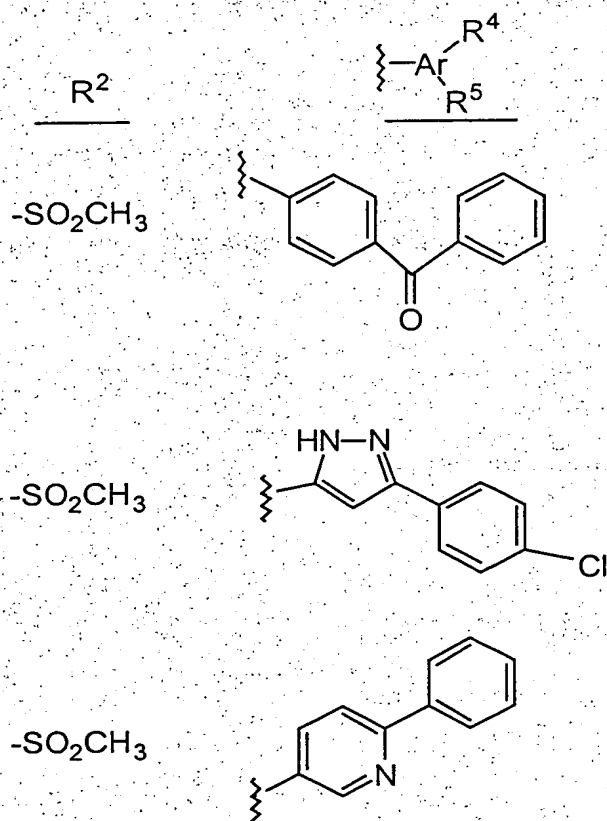
or a pharmaceutically acceptable salt thereof.

- 5 13. The compound of Claim 12 wherein X is N, Z is -CH= and V and W are both -CH=.

14. The compound of Claim 13, or a pharmaceutically acceptable salt thereof selected from those depicted in the following Table

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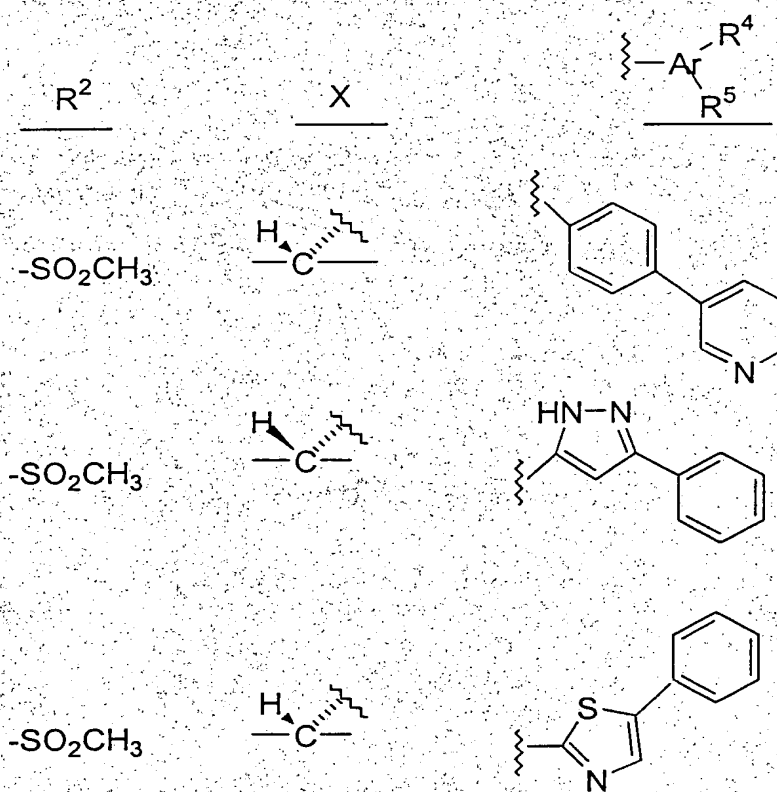


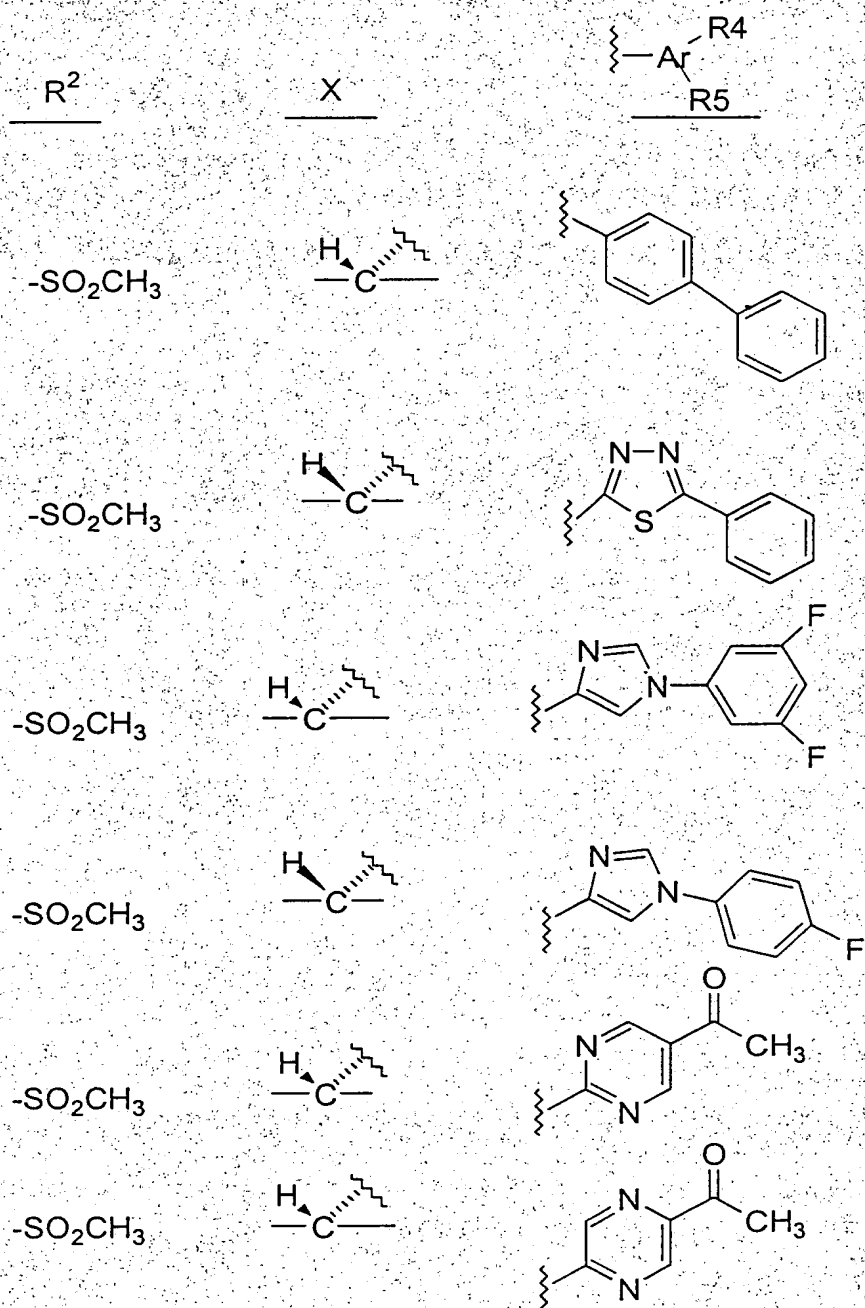


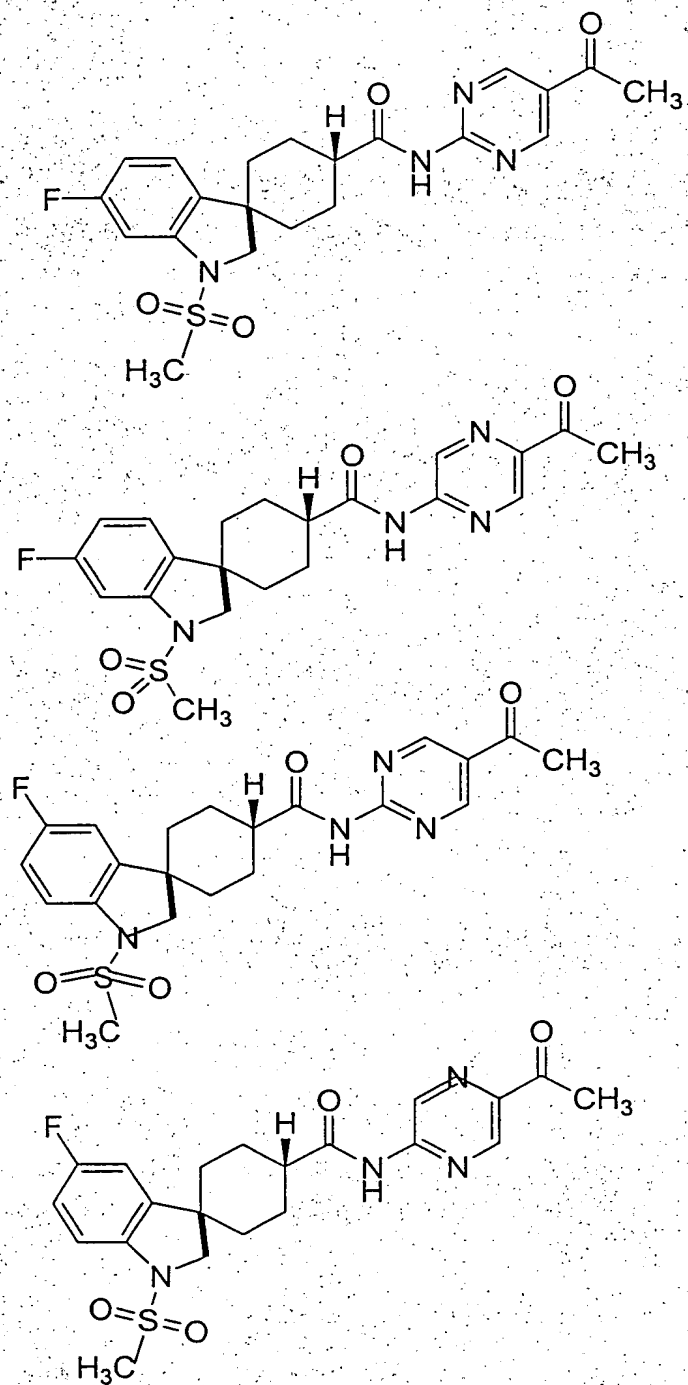
15. The compound of Claim 12 wherein X is -CH=, Z is N and V and W are both -CH=.

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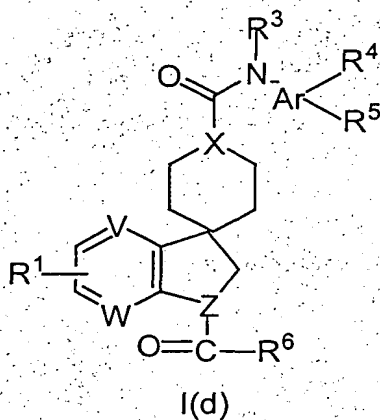
16. The compound of Claim 15 or a pharmaceutically acceptable salt thereof, selected from those depicted in the following Table;





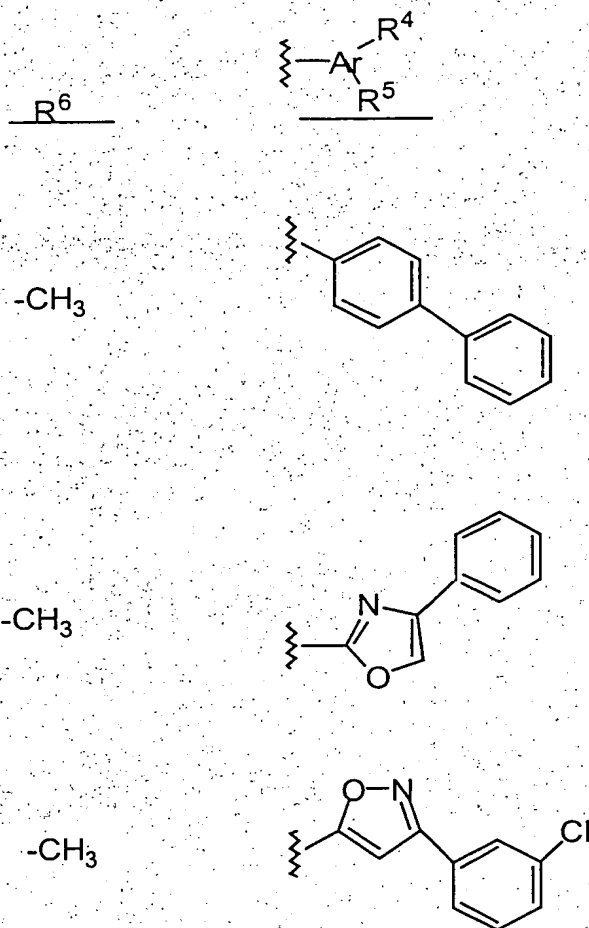


17. The compound of Claim 1 wherein R² is -COR⁶ of structural formula I(d):



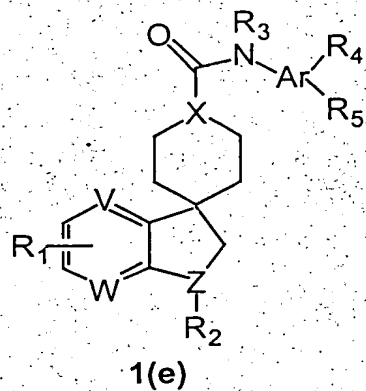
5 or a pharmaceutically acceptable salt thereof.

18. The compound of Claim 17 or a pharmaceutically acceptable salt thereof selected from those depicted in the following Table:

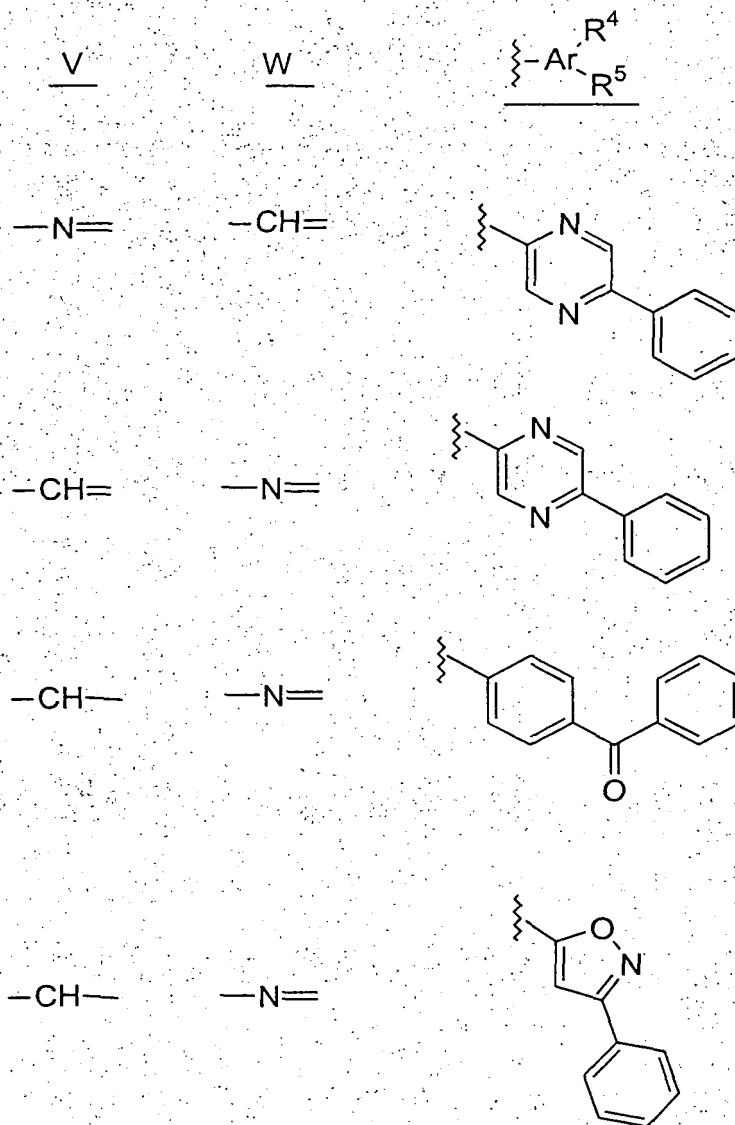


19. The compound of Claim 1 of structural formula I(e), wherein one of V or W is nitrogen and the other is ---CH= .

5



20. The compound of Claim 19 wherein R¹ and R³ are both
hydrogen.
21. The compound of Claim 20 wherein R² is -SO₂CH₃ or -
SO₂NH₂.
22. The compound of Claim 21 selected from the compounds
depicted in the following TABLE



23. A method of treating Y5 receptor mediated diseases which comprises administering to a patient in need of such treatment a non-toxic
 5 therapeutically effective amount of a compound of Claim 1 that selectively antagonizes the Y5 receptor in preference to the other NPY receptors.

24. The method of Claim 23 wherein the Y5 mediated disease is obesity.

25. A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and an effective amount of a selective Y5 antagonist.
- 5